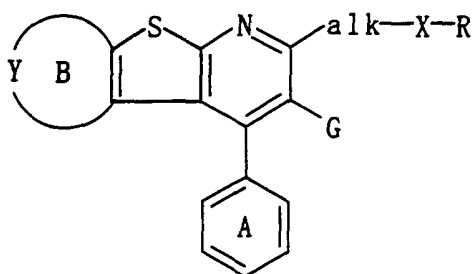


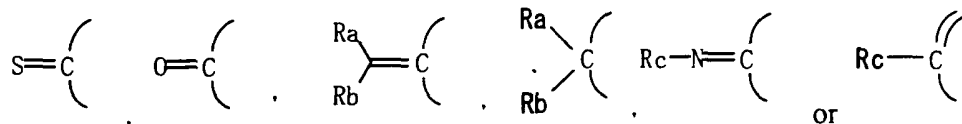
CLAIMS

1. A compound represented by the formula (I):



(I)

5 wherein G represents a halogen atom, hydroxyl group, an optionally substituted amino group, an optionally substituted lower alkyl group or an optionally substituted alkoxy group; alk represents an optionally substituted lower alkylene group; X represents oxygen atom, an
 10 optionally oxidized sulfur atom, or $-(CH_2)_q-$ (q represents an integer of 0 to 5); R represents an optionally substituted amino group or an optionally substituted heterocyclic group; ring B represents an optionally substituted Y-containing 5- to 8-membered ring whose ring
 15 constituent atoms contain no nitrogen atom; Y represents oxygen atom, an optionally oxidized sulfur atom,



(wherein Ra and Rb are the same or different and,

respectively, represent hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group, or Ra and Rb may be combined each other to form a 5- to 7-membered ring; and Rc represents hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group); and ring A represents an optionally substituted benzene ring, or a salt thereof.

2. The compound according to claim 1, wherein alk is methylene.

3. The compound according to claim 1, wherein G is a halogen atom.

4. The compound according to claim 1, wherein G

is chlorine atom.

5. The compound according to claim 1, wherein X is $-(CH_2)_q-$ (q represents an integer of 0 to 5).

6. The compound according to claim 1, wherein X
5 is a bond.

7. The compound according to claim 1, wherein the optionally substituted amino group represented by R is $-N(R^1)(R^2)$ (wherein R^1 and R^2 , which may be the same or different, respectively, represent hydrogen atom, an
10 optionally substituted hydrocarbon group, or an optionally substituted acyl, sulfonyl, sulfinyl, or heterocyclic group, or R^1 and R^2 may be combined each other to form an optionally substituted nitrogen-containing 5- to 7-membered heterocyclic group).

8. The compound according to claim 7, wherein
15 R^1 and R^2 are combined each other to form a nitrogen-containing optionally substituted 5- to 7-membered heterocyclic group.

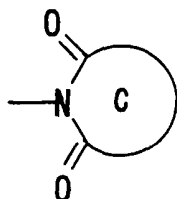
9. The compound according to claim 7, wherein
20 R^1 and R^2 are acyl groups.

10 The compound according to claim 1, wherein R is an optionally substituted nitrogen-containing heterocyclic group.

11. The compound according to claim 1, wherein
25 the substituent on the optionally substituted heterocyclic

group represented by R is oxo group.

12. The compound according to claim 1, wherein R is

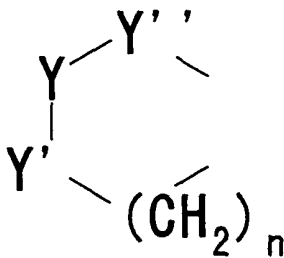


5 wherein ring C represents a 5- to 7-membered heterocyclic group optionally containing one or more hetero atoms selected from nitrogen, sulfur and oxygen atoms, in addition to the nitrogen atom.

13. The compound according to claim 1, wherein
10 ring B is an optionally substituted 6-membered ring containing Y.

14. The compound according to claim 1, wherein the substituents on ring B are one to four substituents selected from a C_{1-6} alkyl group and a halogen atom.

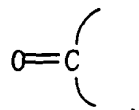
15 15. The compound according to claim 1, wherein ring B is a ring represented by the formula:



wherein Y' and Y'' represent carbon atom, sulfur atom or

oxygen atom, respectively; n represents an integer of 0 to 4; and Y is as defined in claim 1.

16. The compound according to claim 1, wherein Y is an optionally oxidized sulfur atom or



5

17. The compound according to claim 1, wherein ring A is benzene ring which may be substituted with one to four substituents selected from a halogen atom, nitro group, an optionally substituted alkyl group, an optionally substituted hydroxyl, an optionally substituted thiol group, an optionally substituted amino group, an optionally substituted acyl group, an optionally esterified carboxyl group, and an optionally substituted aromatic ring group.

18 The compound according to claim 1, wherein the substituent on ring A is a C₁₋₆ alkoxy group or hydroxyl group.

19 The compound according to claim 1, wherein the compound represented by formula (I) is:

1-{{3-chloro-4-(4-methoxyphenyl)-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl}methly}-2,5-pyrrolidinedione ethylene ketal;
3-{{3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl}methyl}-1,3-thiazolidine-2,4-dione or an optically active compound

20

thereof or a salt thereof;

3-{[3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl}-1,3-oxazolidine-2,4-dione or an optically active compound

5 thereof or a salt thereof;

1-{[3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl}-2,5-pyrrolidinedione or an optically active compound thereof or a salt thereof;

10 1-{[3-chloro-4-(4-methoxyphenyl)-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl}-2,5-pyrrolidinedione or a salt thereof;

1-[[3-chloro-4-(4-methoxyphenyl)-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl]-2,5-pyrrolidinedione or a salt thereof;

3-{[3-chloro-4-(4-methoxyphenyl)-8,8-dimethyl-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl}-1,3-thiazolidine-2,4-dione or a salt thereof;

1-[[3-chloro-4-(4-methoxyphenyl)-6,6-dimethyl-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl]-2,5-pyrrolidinedione or a salt thereof; or
2-aminomethyl-3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridine or an optically active compound thereof or a salt thereof.

25

20. The compound according to claim 1, wherein

the compound represented by formula (I) is:

3-([3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl)-1,3-thiazolidine-2,4-dione or an optically active compound thereof or a salt thereof;

3-([3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl)-1,3-oxazolidine-2,4-dione or an optically active compound thereof or a salt thereof; or

1-([3-chloro-4-(4-methoxyphenyl)-7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl)-2,5-pyrrolidinedione or an optically active compound thereof or a salt thereof.

21. The compound according to claim 1, wherein the compound represented by formula (I) is:

1-([3-chloro-4-(4-methoxyphenyl)-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl)-2,5-pyrrolidinedione or a salt thereof;

1-([3-chloro-4-(4-methoxyphenyl)-8,8-dimethyl-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl)-2,5-pyrrolidinedione or a salt thereof;

3-([3-chloro-4-(4-methoxyphenyl)-8,8-dimethyl-7-oxo-5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl)-1,3-thiazolidine-2,4-dione or a salt thereof; or

1-([3-chloro-4-(4-methoxyphenyl)-6,6-dimethyl-7-oxo-

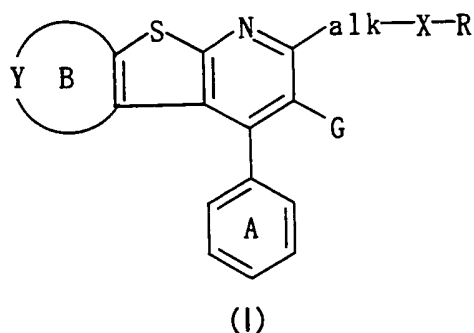
5,6,7,8-tetrahydro[1]benzothieno[2,3-b]pyridin-2-yl)methyl)-2,5-pyrrolidinedione or a salt thereof.

22. The compound according to claim 1, wherein the compound represented by formula (I) is:

5 1-{[3-chloro-4-(4-hydroxyphenyl)-7,7-dioxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-2-yl)methyl]-2,5-pyrrolidinedione or a salt thereof;
4-{3-chloro-2-[(2,5-dioxo-1-pyrrolidinyl)methyl]-7,7-oxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-4-
10 yl)phenyl} diisobutyl phosphate or a salt thereof; or
butyl 4-{3-chloro-2-[(2,5-dioxo-1-pyrrolidinyl)methyl]-7,7-dioxido-5,8-dihydro-6H-thiopyrano[4',3':4,5]thieno[2,3-b]pyridin-4-yl)phenyl carbonate or a salt thereof.

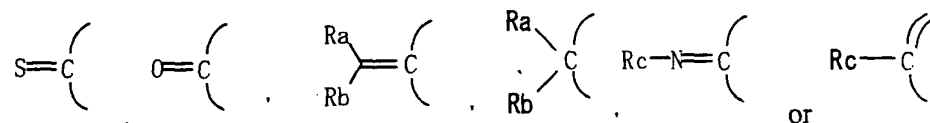
23. A prodrug of the compound according to claim
15 1.

24. A process for producing a compound represented by the general formula (I):



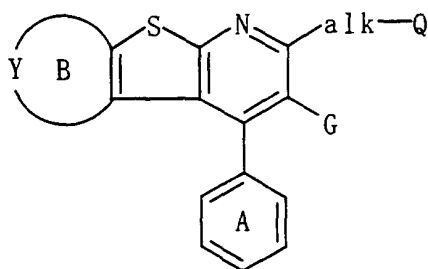
wherein G represents a halogen atom, hydroxyl group, an
20 optionally substituted amino group, an optionally

substituted lower alkyl group or an optionally substituted alkoxy group; alk represents an optionally substituted lower alkylene group; X represents oxygen atom, an optionally oxidized sulfur atom, or $-(CH_2)_q-$ (q represents an integer of 0 to 5); R represents an optionally substituted amino group or an optionally substituted heterocyclic group; ring B represents an optionally substituted Y-containing 5- to 8-membered ring whose ring constituent atoms contain no nitrogen atom; Y represents oxygen atom, an optionally oxidized sulfur atom,



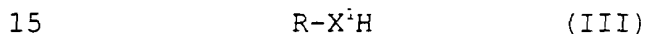
(wherein Ra and Rb are the same or different and, respectively, represent hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group, or Ra and Rb may be combined each other to form a 5- to 7-membered ring; and Rc represents hydrogen atom, a halogen atom, an optionally

substituted hydrocarbon group, an optionally substituted
acyl group, an optionally substituted carbamoyl group, an
optionally substituted thiocarbamoyl group, an optionally
substituted sulfonyl group, an optionally substituted
5 sulfinyl group, an optionally substituted hydroxyl group,
an optionally substituted thiol group, an optionally
esterified carboxyl group, or an optionally substituted
heterocyclic group); and ring A represents an optionally
substituted benzene ring, or a salt thereof which comprises
10 reacting a compound represented by the formula (II-1):

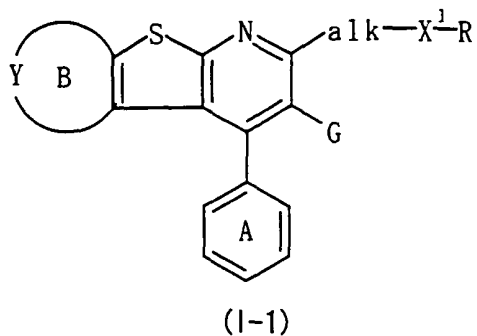


(II-1)

wherein Q represents a leaving group; and the other symbols
are as defined above, or a salt thereof, with a compound
represented by the formula (III):



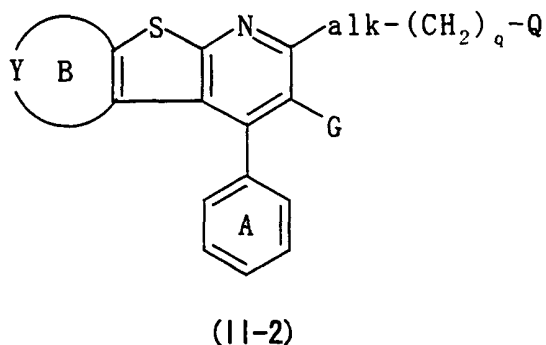
wherein R is as defined above; and X^i represents oxygen
atom, or an optionally oxidized sulfur atom, or a salt
thereof to obtain a compound represented by the formula (I-
1):



wherein each symbol is as defined above, or a salt thereof;

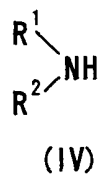
or

5 reacting a compound represented by the formula
(II-2):



wherein each symbol is as defined above, or a salt thereof,

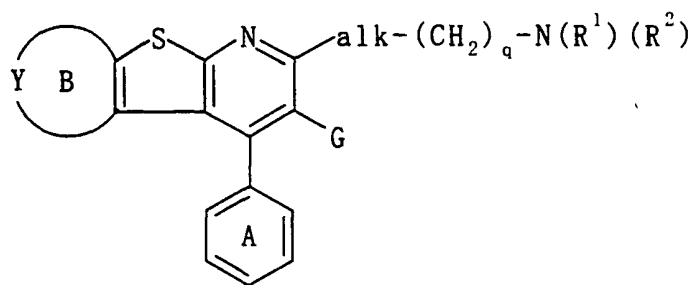
with a compound represented by the formula (IV):



10

wherein R^1 and R^2 , which may be the same or different, respectively, represent an optionally substituted hydrocarbon group, an optionally substituted acyl group, an

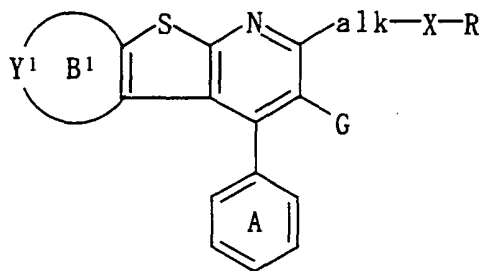
optionally substituted sulfonyl group, or an optionally substituted heterocyclic group, or R^1 and R^2 may be combined each other to form an optionally substituted nitrogen-containing 5- to 7-membered ring, or a salt thereof to
 5 obtain a compound represented by the formula (I-2):



(I-2)

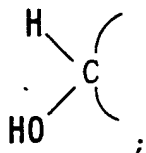
wherein each symbol is as defined above, or a salt thereof;
 or

subjecting a compound represented by the formula
 10 (I-3):

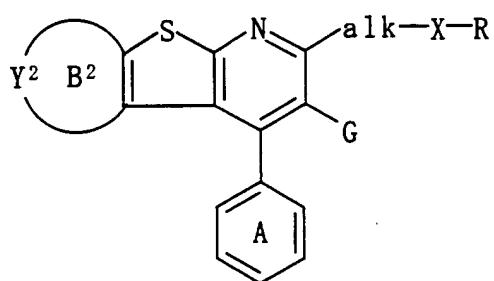


(I-3)

wherein ring B^1 represents an optionally substituted Y^1 -containing 5- to 8-membered ring whose ring constituent atoms contain no nitrogen atom; Y^1 represents sulfur atom
 15 or

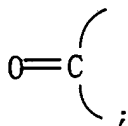


and the other symbols are as defined above, or a salt thereof to oxidation to obtain a compound represented by the formula (I-4):



(I-4)

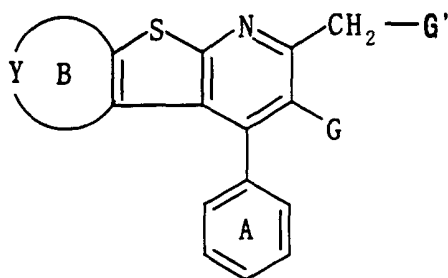
wherein ring B² represents an optionally substituted Y²-containing 5- to 8-membered ring whose constituent ring atoms contain no nitrogen atom; Y² represents an oxidized sulfur atom or



and the other symbols are as defined above, or a salt thereof; or

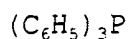
reacting a compound represented by the formula

(II-3):

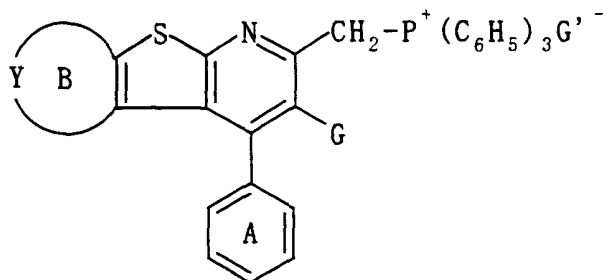


(II-3)

wherein G' represents a halogen atom; and the other symbols are as defined above, or a salt thereof, with

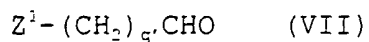


5 in a solvent to obtain a compound represented by formula (VI):

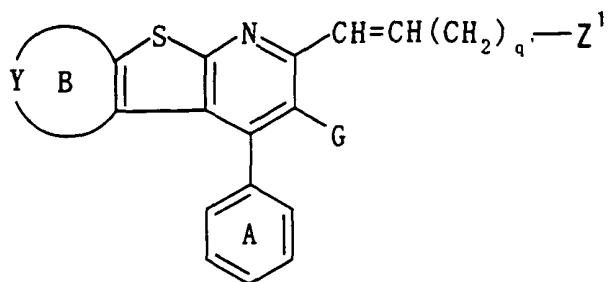


(VI)

wherein each symbol is as defined above, then reacting the compound represented by the formula (VI) with a compound
10 represented by the formula (VII):

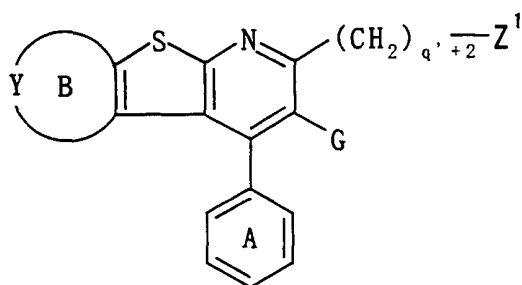


wherein Z^1 represents an optionally substituted heterocyclic group; and q' represents an integer of 0 to 4, or a salt thereof to obtain a compound represented by
15 formula (VIII):



(VIII)

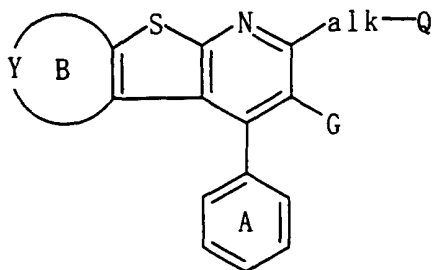
wherein each symbol is as defined above, or a salt thereof,
 and further subjecting the compound represented by formula
 (VIII) or a salt thereof to reduction to obtain a compound
 5 represented by the formula (I-5):



(I-5)

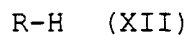
wherein each symbol is as defined above, or a salt thereof;
 or

reacting a compound represented by the formula
 10 (II-1):

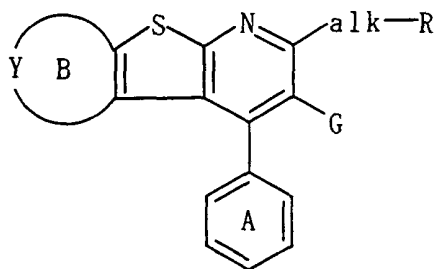


(II-1)

wherein each symbol is as defined above, or a salt thereof,
with a compound represented by formula (XII):



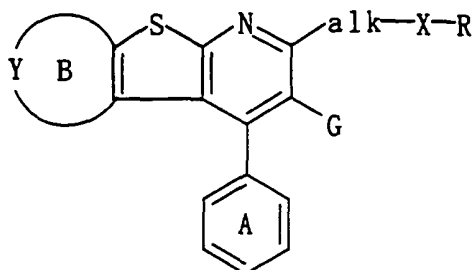
- 5 wherein R is as defined above, or a salt thereof to obtain
a compound represented by the formula (I-9):



(I-9)

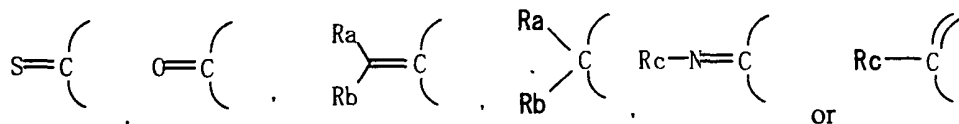
wherein each symbols is as defined above, or a salt thereof.

25. A pharmaceutical composition comprising a
10 compound represented by the general formula (I):



(I)

wherein G represents a halogen atom, hydroxyl group, an optionally substituted amino group, an optionally substituted lower alkyl group or an optionally substituted alkoxy group; alk represents an optionally substituted lower alkylene group; X represents oxygen atom, an optionally oxidized sulfur atom, or $-(CH_2)_q-$ (q represents an integer of 0 to 5); R represents an optionally substituted amino group or an optionally substituted heterocyclic group; ring B represents an optionally substituted Y-containing 5- to 8-membered ring whose ring constituent atoms contain no nitrogen atom; Y represents oxygen atom, an optionally oxidized sulfur atom,



(wherein Ra and Rb are the same or different and, respectively, represent hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl

group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group, or Ra and Rb may be combined each other to form a 5- to 7-membered ring; and Rc represents hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group); and ring A represents an optionally substituted benzene ring, a prodrug thereof, or a pharmaceutically acceptable salt thereof.

26. The pharmaceutical composition according to claim 25 for prevention or treatment of inflammatory diseases.

27. The pharmaceutical composition according to claim 25 for prevention or treatment of arthritis.

28. The pharmaceutical composition according to claim 25 for prevention or treatment of rheumatism.

29. The pharmaceutical composition according to claim 25 for prevention or treatment of chronic rheumatoid arthritis.

5 30. The pharmaceutical composition according to claim 25, which is a bone resorption suppressant.

31. The pharmaceutical composition according to claim 25 for prevention or treatment of osteoporosis.

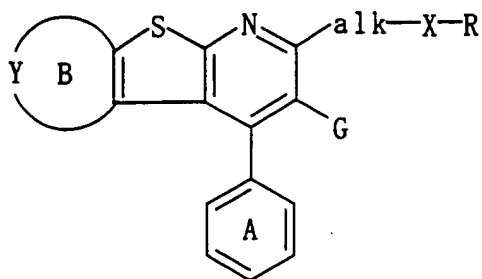
32. The pharmaceutical composition according to claim 25, which is a suppressant of cytokine production.

10 33. The pharmaceutical composition according to claim 25 for prevention or treatment of autoimmune diseases.

34. The pharmaceutical composition according to claim 25 for prevention or treatment of rejection reaction after organ transplantation.

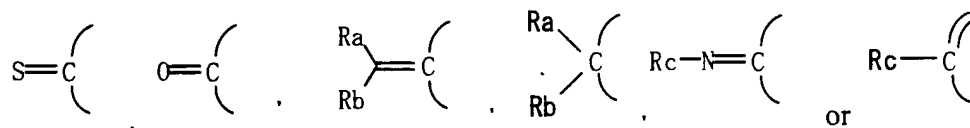
15 35. The pharmaceutical composition according to claim 25, which is a T-cell differentiation-controlling medicament.

36. A method for preventing or treating inflammatory diseases which comprises administering an
20 effective amount of a compound represented by the formula (I):



(I)

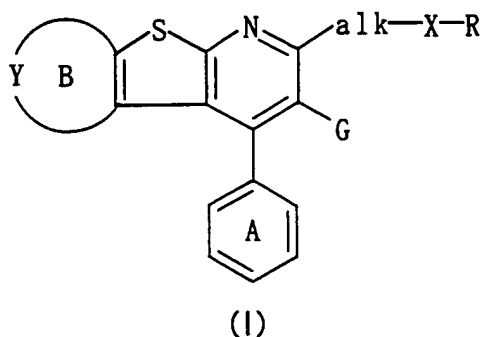
wherein G represents a halogen atom, hydroxyl group, an optionally substituted amino group, an optionally substituted lower alkyl group or an optionally substituted alkoxy group; alk represents an optionally substituted lower alkylene group; X represents oxygen atom, an optionally oxidized sulfur atom, or $-(CH_2)_q-$ (q represents an integer of 0 to 5); R represents an optionally substituted amino group or an optionally substituted heterocyclic group; ring B represents an optionally substituted Y-containing 5- to 8-membered ring whose ring constituent atoms contain no nitrogen atom; Y represents oxygen atom, an optionally oxidized sulfur atom,



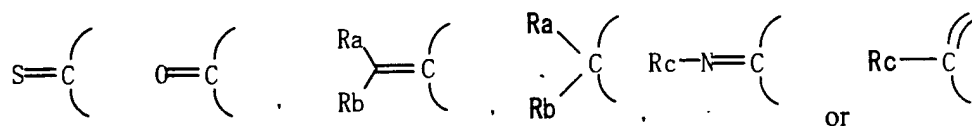
(wherein Ra and Rb are the same or different and, respectively, represent hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl

group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group, or Ra and Rb may be combined each other to form a 5- to 7-membered ring; and Rc represents hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group); and ring A represents an optionally substituted benzene ring, a prodrug thereof, or a pharmaceutically acceptable salt thereof, to a mammal in need of the prevention or treatment.

37. Use of a compound represented by the formula (I):



wherein G represents a halogen atom, hydroxyl group, an optionally substituted amino group, an optionally substituted lower alkyl group or an optionally substituted alkoxy group; alk represents an optionally substituted lower alkylene group; X represents oxygen atom, an optionally oxidized sulfur atom, or $-(CH_2)_q-$ (q represents an integer of 0 to 5); R represents an optionally substituted amino group or an optionally substituted heterocyclic group; ring B represents an optionally substituted Y-containing 5- to 8-membered ring whose ring constituent atoms contain no nitrogen atom; Y represents oxygen atom, an optionally oxidized sulfur atom,



(wherein Ra and Rb are the same or different and, respectively, represent hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl

group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group, or Ra and Rb may be combined each other to form a 5- to 7-membered ring; and Rc represents hydrogen atom, a halogen atom, an optionally substituted hydrocarbon group, an optionally substituted acyl group, an optionally substituted carbamoyl group, an optionally substituted thiocarbamoyl group, an optionally substituted sulfonyl group, an optionally substituted sulfinyl group, an optionally substituted hydroxyl group, an optionally substituted thiol group, an optionally esterified carboxyl group, or an optionally substituted heterocyclic group); and ring A represents an optionally substituted benzene ring, a prodrug thereof, or a pharmaceutically acceptable salt thereof in the manufacture of a pharmaceutical composition for preventing or treating inflammatory diseases.